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Two new species of myxosporidians, Myxosoma channai n.sp. and Myxobolus tripathii n.sp. from fresh water fishes of Andhra Pradesh

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Abstract. Two new species of myxosporidians, Myxosoma channai n.sp. infecting the fins, body muscles, liver and kidney of Channa punctata Bl. and Myxobolus tripathii n.sp. infecting the gut and the liver of Clarius sp. are described. A checklist of all the species of myxosporidians reported from fishes of India is also given.

Keywords. Myxosoma channai; Myxobolus tripathii n.sp.; Channa punctata; Clarius sp.; new species.

1. Introduction

Southwell and Prashad (1918) reported the first myxosporidian from Indian fishes. Later Ray (1933), Chakravarty (1939, 1943), Ganapati (1941), Setna (1942), Chakravarty and Basu (1948), Tripathi (1952), Qadri (1962a-d, 1965, 1967, 1969 and 1970), Bhatt and Siddiqui (1964), Qadri and Lalitakumari (1965), Lalitakumari (1969), Chaudhuri and Chakravarty (1970), Choudhury and Nandi (1973), Narasimhamurti (1970), Narasimhamurti and Kalavati (1975, 1979a-c), Narasimhamurti et al (1980) described a number of myxosporidians belonging to different genera from fishes of India. Tripathi (1952) gave a review of the work done till 1952 and gave a checklist of all the myxosporidians reported till that time.

So far about 80 species of myxosporidia (table 2) have been described from Indian fishes. A checklist of all the species of myxosporidians described so far from India is included in the present paper because there is no such compilation after 1952 (Tripathi 1952).

While examining the fresh water fishes of Visakhapatnam District we came across 2 myxosporidians, one belonging to the genus *Myxosoma* infecting the fins, body muscles, liver and kidney of *Channa punctata* B1. and the other belonging to the genus *Myxobolus* infecting the muscles of the gut and liver of *Clarius* sp. Both of them are considered new to science and are described in this paper.

2. Materials and methods

Channa punctata B1. were collected from 3 different places in Visakhapatnam District; from a fresh water tank used mostly for drinking purposes located at the foot of hill in Srungavarapukota, about 30 miles north of Visakhapatnam; from an abandoned tank which has highly polluted water and plenty of green algae in Elamanchili, about 38 miles south of Visakhapatnam and a stream near the Dairy farm in Visakhapatnam. Clarius sp. were collected from a tank near the Visakhapatnam Port.

Different parts of the body of the fish were examined for myxosporidian parasites and when infection was detected as evident by the presence of cysts, smears were prepared, air-dried and fixed in methyl alcohol, hydrolysed in 1N HCl at 60°C for 10 min and stained with Giemsa. Smears were also wet-fixed in Schaudinn's or Carnoy's fluid and stained with Heidenhain's iron haematoxylin or according to Feulgen's technique. Fresh spores were treated with india ink and lugol's iodine to detect the presence of any mucous envelope and iodinophilous vacuole. The measurements and drawings of the spores were made in the fresh condition.

3. Observations

Myxosoma channai n. sp. Host: Channa punctata Bl.

Site of infection: Fins, body muscles, kidney and liver.

Type slides: Author's Collection at Department of Zoology, Andhra University, Waltair (Andhra Pradesh).

68 out of 132 (51.5%) of Channa punctata Bl. collected from 3 different localities in Visakhapatnam District during February-April 1979 were, found infected with a new species of myxosoma. The size of the fish, the site and percentage of infection differed in fish collected in the different places (table 1).

32 fish brought from Srungavarapukota were maintained in aquaria tanks and fed with commercially available fish food. Infected fish from this area showed infection predominantly in the kidney and to a lesser extent in the liver. The

Table 1. Incidence of M. Channai in fishes collected from different localities in Visakhapatnam District.

| Place | Site of infection | Size | Number examined | Number infected |
|------------------------------------|-------------------|---------------------------------|--------------------|--------------------|
| Fresh water tank Srungavarapukota | Kidney and Liver | 6″-7″ | 32 | 24 |
| Fresh water tank Elamanchili | Body muscles | 4″-6′′ | 36 | 4 |
| Dairy farm stream Visakhapatnam | Fins | $2\frac{1}{2}''-3\frac{1}{2}''$ | 32 | 20 |

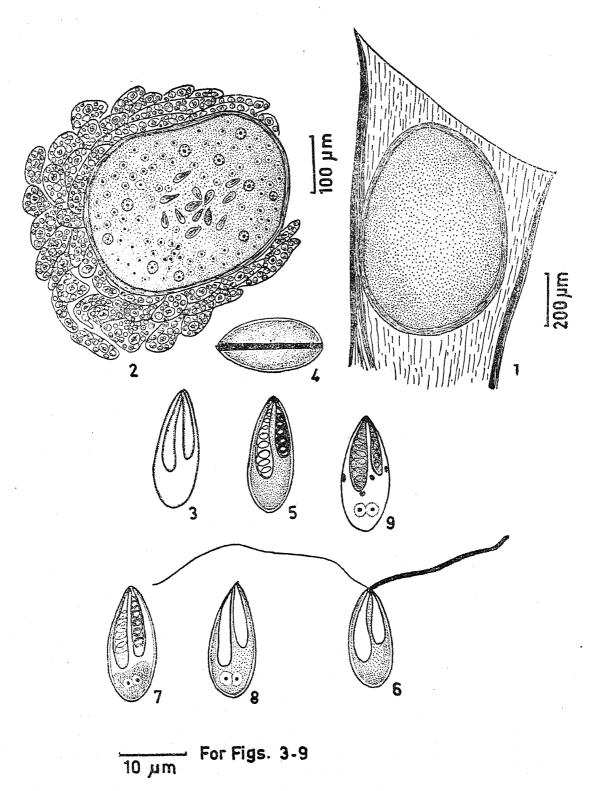
infection was heavy and each fish yielded about 10 cysts. Five out of the 7 fish which were sacrificed immediately after being brought to the laboratory showed heavy infection. More cysts were either superficially attached or deeply embedded in the kidneys and a lesser number in the liver. Unfortunately all the cysts were immature and did not show any spores. The rest of the 25 fish were maintained in the laboratory. Five of them died in 7 days and they showed infection both in the kidneys and the liver, heavier in the former than in the latter. Sections of the kidney showed cysts with few spores in the centre and developmental stages along the periphery (figure 2). The remaining 20 fish were maintained in the laboratory for another 2 weeks. The fish appeared active and there were no casualities during that period. All the fish were sacrificed at the end of 2 weeks and examined for infection and 14 of them showed heavy infection in the kidneys.

Four of the 36 fish (11%) brought from Elamanchili showed the infection limited to the body muscles. The size of the cysts varied from 0.2-0.5 mm in diameter. The infection was lighter and yielded 3 and 4 cysts in each of the 2 infected fish.

Twenty out of the 32 fish collected from the stream near the dairy farm area, Visakhapatnam showed infection limited to the fins. The infection was heavy and as many as 10 cysts were recovered from each fish. The infection was prevalent in the smaller fish and was absent in the larger fish. Cysts measuring $800-1020~\mu m$ in diameter were either spherical or oval and were surrounded by a thin membrane (figure 1).

Sections of early cysts showed two types of nuclei, the larger nuclei which were fewer in number had a conspicuous large endosome and a distinct nuclear membrane and were probably the generative nuclei while the smaller ones which were more numerous had a small dot-like endosome and an indistinct nuclear membrane which were probably the somatic nuclei (figure 2).

Spores: The spores were pale brown in colour, elongately oval and had a bluntly pointed anterior end and a rounded posterior end. They measured 14.5-18 x $6-6.5 \mu m$. The two spore valves meet along a conspicuously thick median sutural ridge (figure 4). Two polar capsules which are elongate and pear-shaped were of dissimilar size and were situated one on either side of the median line. The larger polar capsule measured $9.0-10.8 \times 2.8-3.2 \,\mu m$ and the smaller one measured $7.2-8.8 \times 2.8-3.2$ (figure 5). The polar filament in the larger polar capsule was thin with 8 coils wound in an anticlock wise direction and when fully everted measured 35-42 μ m in length. The polar filament from the smaller polar capsule was significantly thick, deeply stained and appeared almost solid and had 5-6 coils wound in a clock-wise direction in the polar capsule. When fully everted the polar filament measured $20-28 \mu m$ in length. The sporoplasm was triangular in shape with rounded corners and the base was directed anteriorly (figure 7). Two deeply stained nuclei were placed a little apart along the longitudinal axis. The endosome was deeply stained but the nuclear membrane was not clear (figure 8). Some of the developing sporoblasts showed 6 nuclei, 2 lying below the capsule and 2 near the valves (between the polar capsule and the spore wall) which showed fine deeply stained chromatin granules dispersed over a wide area. A distinct nuclear membrane was not observed. The other 2 nuclei were located in the sporoplasm and have a distinct endosome and a delicate nuclear membrane.



Figures 1-9. Myxosoma channai n.sp. 1. A cyst attached to the fin. 2. T.S. kidney showing early cyst. 3. A fresh spore. 4. Sutural view of the spore. 5. A spore stained with Giemsa. 6. Spore showing extruded polar filaments: Note polar filaments of unequal thickness and length. 7. Spores stained with iron haematoxylin. Note binucleate sporoplasm. 8. Spore stained with Feulgen. 9. A developing sporoblast.

4. Discussion

16 species of myxosporidians belonging to 5 genera, Myxobolus, Unicauda, Henneguya, Zschokella and Myxosoma are reported from 3 species of Channa (=Ophice-phalus), C. punctata, C. striata and C. gachua from different parts of India (table 2).

Many species of Myxosoma have been reported from a variety of marine and fresh water fishes from all over the world and 6 of them are from India, 3 of them from fresh water fishes of Hyderabad (Lalitakumari 1969) and 3 are from estuarine fishes of Waltair (Narasimhamurti 1970; Narasimhamurti and Kalavati 1979). Lalitakumari (1969) reported 3 species of Myxosoma, M. indiae, (12·4-15·0 × $6\cdot4-8\cdot6~\mu\text{m}$), M. hyderabadense $(9\cdot3-11\cdot5\times5\cdot0-8\cdot0~\mu\text{m})$ and M. andhrae $(12\cdot1 15.7 \times 5.7 - 8.6 \,\mu\mathrm{m}$). Narasimhamurti (1970) and Narasimhamurti and Kalavati (1979) reported 3 more species, M. intestinalis ($12 \cdot 5 - 13 \cdot 5 \times 8 \cdot 6 - 9 \cdot 5 \mu m$), M. lairdi $(9.0-9.5 \times 4.6-5.3 \,\mu\text{m})$ and M. microspora $(4.8-5.2 \,\mu\text{m})$ in diameter). Among them the spores of M. hyderabadense, M. lairdi and M. microspora are much smaller than the present form (14.5–18.0 \times 6.0–6.5 μ m) though the polar capsules were of a similar size. The spores of M. indiae, M. andhrae and M. intestinalis are bigger in size but are still smaller than the present form. The present form resembles M. indiae and M. andhrae in having unequal sized polar capsules. In M. indiae the unequal polar capsules were stated to be 2 or 3 in number and are smaller than those in the present form. The sporoplasm in M. indiae extends a little into the intercapsular space and hence differs from the present form. The spores of M. andhrae (which has the same host as the present form) though large are smaller than the present form and posses parietal folds which are absent in the present form. The present form resembles M. andhrae in having unequal-sized polar capsules but differs from it because the sporoplasm extends into the intercapsular space upto the anterior pole while in the present case it is triangular and is located at the posterior pole. Some of the spores in M. andhrae show an appendage measuring $1.0-3.6 \mu m$ which is absent in the present form. The polar filaments in M. andhrae appear to be similar (as seen from figure 20) while they appear and measure differently in the present form. The cysts in M. andhrae were attached to the gut by means of small fibres while in the present case the cysts are attached to fins or tissues directly or embedded in the tissue. Thus the only species of Myxosoma, M. andhrae which is reported from the same host as the present form differs considerably, hence this is considered a new species and the name Myxosoma channai n. sp. after the host is proposed for the same.

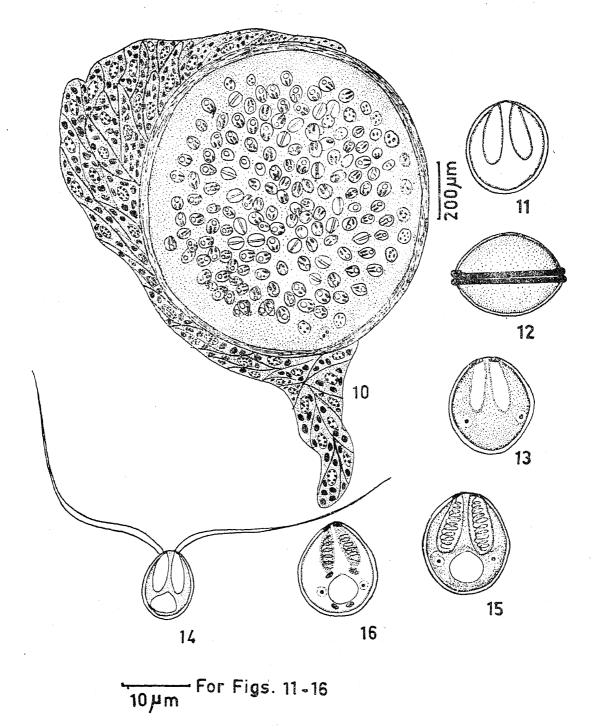
Myxobolus tripathii n. sp. Host: Clarius sp.

Site of infection: Wall of gut and visceral organs.

Type slides: Author's collection and Department of Zoology, Andhra University,

Waltair (Andhra Pradesh).

3 out of 75 Clarius sp. ranging in size from 7.5-8.0 inches and collected from a tank near Visakhapatnam Port were found infected with a new species of Myxo-bolus. Opaque white cysts ranging in diameter from 0.5-1.0 mm were either superficially attached to the gut epithelium or deeply embedded in the liver tissue. The cyst wall was thin and single. Cellular hypertrophy associated with vacuo-



Figures 10-16. Myxobolus tripathii n.sp. 10. T.S. liver showing an attached cyst. 11. A fresh spore. 12. Sutural view of the spore. 13. A spore treated according to Feulgen's technique. 14. Spore showing extruded polar filaments. 15. Spore stained with Giemsa: Note coiled polar filament. 16. A developing sporoblast.

lation of the cytoplasm was observed in the adjacent tissue cells. In some of the cells the nucleus also showed hypertrophy but there was no rupture of the cell (figure 10).

Fresh spores were either rounded or oval measuring $9.8 - 10.2 \times 12.0 - 13.5 \,\mu\mathrm{m}$ (figure 11). The spore walls were symmetrical and meet along the thickened sutural ridge which was further thickened at the poles (figure 12). 2 oval polar capsules of equal size measuring $5.0-6.0 \times 2.5 \,\mu\mathrm{m}$ were present, one on either side of the median line and they open independently to the outside. A prominent cushion-like thickening which was strongly basophilic was present at the openings (figure 13). The polar filaments showed 8 coils in each of the polar capsules and they were coiled in an anticlock wise direction. The polar filaments were uniformly thin and when fully everted measured 45-50 μm in length (figure 14). An oval iodinophilous vacuole measuring $3 \cdot 6 - 4 \cdot 5 \,\mu\mathrm{m}$ which was stained with Lugol's iodine, positive to PAS and Best's carmine and negative to PAS after saliva digestion was present at the posterior pole (Figures 15 and 16). Similar tests conducted on the spores of the new species of Myxosoma reported earlier in this paper were negative. The sporoplasm was binucleate and extended like a rim round the iodine vacuole. 2 widely separated vesicular nuclei were present in the sporoplasm (figure 15). In immature spores 2 capsulogenous nuclei at the base of the polar capsules and 2 valvulogenous nuclei at the posterior end below the iodine vacuole were usually present in addition to the 2 sporoplasmic nuclei situated on either side of the iodinophilous vacuole (figure 16).

4. Discussion

The present form is placed in the genus Myxobolus because of the presence of an iodinophilous vacuole in the spore (Kudo 1920). Walliker (1968) synonymised the genus Myxosoma and Myxobolus because of the variability of occurrence of iodine vacuole. Lom (1969) endorsed the view of Walliker, thus abolishing the family Myxosomatidae. However, Galinsky and Meglitsch (1969) felt that the vacuole is useful and a well established taxonomic character. Podlipaivi (1974) stated that the iodine vacuole is "a real morphological formation in the spores of Myxobolidae and should be used in systematics". Moser and Noble (1977), Wold and Iverson (1978) expressed a similar opinion in the matter. Since a distinct iodine vacuole is present in the present form and accepting the opinion expressed by the above authors that the iodine vacuole has taxonomic importance, we propose to assign the parasite described here to the genus Myxobolus Buetschli 1882.

To date as many as 133 species of Myxobolus have been reported from a variety of marine and fresh water fishes from different parts of the world. Among them 19 species are described from the fresh water fishes of India. A perusal of table 2 shows that the only previous record of Myxobolus from Clarius is M. clarii (Chakravarty 1943) from the gall bladder, liver, testes, ovary and fat bodies. The cysts of the present form are bigger in size when compared to the cysts of M. clarii. The spores in M. clarii measured $11 \cdot 3 - 12 \cdot 4 \times 10 \cdot 3 \mu m$, were subspherical in shape with a distinct straight sutural ridge whereas in the form described here the spores are spherical or ovoidal and have a thick sutural ridge which is further

Table 2. Checklist of myxosporidian parasites reported from the fresh water fishes of India.

| | Parasite | Host | Site of infection | Spore measurements (microns) |
|-------|---|--|--|--|
| | | Ord.: Unipolarina Fam: Ceratomyxidae Genus: Ceratomyxa Thelohan 1892 | | |
| . 2 | C. hilsae Chakravarty 1939 C. gobiodesi Chakravarty 1939 | Hilsa ilisha Odontamblyopus rubicunda ; | Gall bladder Gall bladder | 25–40 i4–15 |
| ų 4. | C. scatophagi Chakravarty 1943 C. sagarica Choudhury and and Nandi 1973 | Colisa fasciatus Scatophagus argus Boleophthalmus boddaerti | Gall bladder Bile | 16–26 (breadth) 31 (breadth) |
| | | Genus: Leptotheca Thelohan 1895 | | |
| 6. 7. | L. latesi Chakravarty 1943L. macronesi Chakravarty 1943L. asymmetrica Lalitakumari1969 | Lates calcarifer Macrones gulio Osteochilus neilli | Gall bladder Gall bladder Gills, Gall bladder and intestine | 10·3-12·4 (diameter) 10-14·4 9·4 (breadth) |
| | | Genus Sphaerospora Thelohan 1892 | | |
| ∞ | S. sp. *Southwell and Prashad 1918 | Barilins barna | Under scales | No details |
| | | Genus: Gyrospora Qadri 1962 | | |
| 6 | G. crucifila Qadri 1962 | Labeo fimbriatus | Gills | $9-10 \times 8-8.5$ |
| | | Fam: Chloromyxidae Thelohan 1892 Genus: Chloromyxum Mingazzini 1890 | | |

| $8\cdot 24 	imes 10\cdot 3$ | 9–10 (diameter) 7.2 (diameter) $5.7-7.1 \times 2.5-3.5$ | | $5.5 - 5.8 \times 7.2$ | 9 (diameter) | 9.4×9.8 | | $12 \cdot 4 - 15 \times 6 \cdot 4 - 8 \cdot 6$ $9 \cdot 3 - 11 \cdot 5 \times 5 - 8$ | $12 \cdot 1 - 15 \cdot 7 \times 5 \cdot 7 - 8 \cdot 6$ | $12.5 - 13.5 \times 8.6 - 9.5$ | $4 \cdot 6 - 5 \cdot 2 \times 9 - 9 \cdot 5$ | 4·8-5·2 (diameter) | $14.5-18\times6-6.5$ |
|--|---|-----------------------------|---------------------------------|--|---|---|---|--|--|---|--|---|
| Gall bladder | Gall bladder Gall bladder Gall bladder | | Muscles and peritoneum in oeso- | Optic lobes | Gut | | Gill filaments Gills | Outer wall of intestine | Gut epithelium | Gut epithelium | Gills | Fins, body muscles, liver and kidney |
| Amphipnous cruchia Heteropneustes fossilis Amblypharyngodon mola | Cirrhina mrigala, C. reba Xenentodon cancila Labeo nigripinnis | Genus: Kudoa Meglitsch 1947 | Strongylura strongylura | Mugil cephalus | Sphyraena jello | Fam: Myxosomatidae Genus: Myxosoma Thelohan 1892 | Barbus sarana Barbus pinnarautus | Ophicephalus punctatus | Mugil waigensis | Liza macrolepis | Mugil cephalus | Channa punctata |
| C. amphipnoui Ray 1933 | C. mrigale Tripathi 1952 C. sp. Tripathi 1952 C. hoarei Lalitakumari 1969 | | K. chilakensis Tripathi 1952 | K. tetraspora Narasimhamurti and Kalavati 1979 | K. sphyraeni Narasimhamurti and Kalavati 1979 | | M. indiae Lalitakumari 1969M. hyderabadense Lalithakumari 1969 | M. andhrae Lalitakumari 1969 | M. intestinalis Narasimhamurti and Kalavati 1970 | M. lairdi Narasimhamurti and Kalavati 1979 | M. microspora Narasimhamurti and Kalavati 1979 | M. channai n. sp. (present record) |
| 10. | 13.5. | | 14. | 15. | 16. | | 17. | 19. | 20. | 77. | 22. | 23. |

* From Kudo (1920).

Table 2. (Contd.)

| | 1 | 2 | 3 | 4 |
|------------|---|---|-------------------------------------|--|
| | | Family: Myxobolidae Genus: Myxobolus Buetschlii 1882 | 23 | |
| 24. 25. | M. mrigale Chakravarty 1939 M. calbausi Chakravarty 1939 | Cirrhina mrigala Labeo calbasu, L. rohita, | Scales Gall bladder | $7.21-8.24 \times 6.18$ $12.4-15 \times 8.2-10$ |
| 26. | M. clarii Chakravarty 1943 | Cirrnna mrigaia Clarius batrachus | Gall bladder, Liver, Testes fat | $11 \cdot 3 - 12 \cdot 4 \times 10 \cdot 3$ |
| 27. | M. cailae Chakravarty 1943 | Catla catla, Labeo rohita, Cirrhina mrioala | Branchiae | $14.5 - 16.5 \times 6.18$ |
| 28. | M. bengalensis Chakravarty | Catla catla | Branchiae | $8.56-9.36 \times 6.42-6.80$ |
| 29. | and basu 1948 M. indicum Tripathi 1952 | Cirrhina mrigala | Scales | $7.21 - 8.24 \times 6.18$ |
| 30. | M. branchialis Tripathi 1952 | Barbus sarana | Gill filaments | $6.4-7 \times 4.5-5 \times 3.2-4$ |
| 31. | | Barbus ticto | Skin | $12.6 - 13.5 \times 9 \times 5.5 - 6.3$ |
| 32. | M. sphericum Tripathi 1952 | Cirrhina mrigala | Scales | y-y-3 × 1.4 × 3-3-3 |
| 33. | M. ophicephali Bhatt and Siddigui 1964 | Ophicephalus punctatus | Accessory respira- rory membrane | $11.6-13.3 \times 4.0-0.3$ |
| 34. | M. aligarhensis Bhatt and Siddioni 1964 | Ophicephalus punctatus | Accessory respiratory membrane | $11 \cdot 4 - 15 \times 6 - 7 \cdot 2$ |
| 35. | M. pinnarauti Lalitakumari 1969 | Barbus pinnarautus | Gill filaments | $7 \times 7 \times 7$ |
| 36. | M. psilorhynchi Lalitakumari 1969 | Psilorhynchus balitora | Gill filaments | $9 \cdot 3 - 10 \cdot 7 \times 8 \cdot 6 - 10$ |
| 37. | M. ampullaceus Lalitakumari 1969 | Barbus kolus | Dorsal and ventral fins | $8.6 - 10.7 \times 6.4 - 7.9$ |
| 38. | M. osmaniae Lalitakumari | Barbus punaubensis | Liver and intestine | $12-15 \times 7 \cdot 1-10$ |
| 39. | M. potaili Lalitakumari 1969 | Labeo potail | Gill contents | $6 \cdot 3 - 7 \cdot 9 \times 4 \cdot 3 - 6 \cdot 4$ |

| Spleen (in all) $12 \cdot 29 - 15 \times 5 \cdot 72 - 7 \cdot 86$ Pharyngeal epi- | thelium (in one) $9.8-12\times 10.2-13.5$ muscles | | Gills $19.8-26.4 \times 5.4-7.2 \times 2-3$ | Mouth cavity $25 \cdot 6 - 39 \cdot 2 \times 3 \cdot 2 - 4 \cdot 4$ Pharyngeal | epithelium $35 \cdot 73 - 45 \cdot 73 \times 14 \cdot 29 - 27 \cdot 15$ | Roof of Buccal $26.4-29.15 \times 11.55-13.75$ cavity | | Gills $26 \cdot 2 - 36 \cdot 2 \times 6 \cdot 3 - 8 \cdot 2$ Branchiae and $41 \cdot 5 - 52 \cdot 6 \times 6 \cdot 18 - 7 \cdot 21$ | Wall of Bulbous $45-52 \times 6-8.5$ | arteriosus Tissues of gill 23.6×2.6 filaments | Gills $52-55 \times 4 \cdot 5-5$ | ntents | appendage) — No measurements Gill filaments $14\cdot6-16\cdot5\times3\cdot2-4$ (Excl. caudal | |
|--|---|---------------------------|---|---|---|---|--------------------------------|--|--------------------------------------|--|--|-----------------------------|--|--|
| Ophicephalus punctatus | Clarius sp. | Genus Unicauda Davis 1944 | Ophicephalus punctatus; Ophicephalus | punctatus | Ophicephalus punctatus | Ophicephalus punctatus | Genus: Henneguya Thelohan 1892 | Lates calcarifer Ophicephalus punctatus | Otolithus ruber, O. maculatus | Ophicephalus punctatus | Notopterus notopterus | | Notopterus notopterus Ophicephalus punctatus | |
| M. punctatus Chaudhuri and Chakravarty 1970 | M. tripathii (present record) | | U. ophicephali Tripathi 1952 | U. basirii Bhatt and Siddiqui1964 | U. bicornuata Chaudhuri and Chakravarty 1970 | U. bengalensis Chaudhuri and Chakravarty 1970 | | H. latesi Chakravarty 1939H. ophicephali Chakravarty1939 | H. otolithi Ganapati 1941 | H. zahoori Bhatt and Siddioui 1964 | H. notopteriae Qadri 1965 H. aadrii I. alitakumari 1065 | H. singhi Lalitakumari 1969 | H. jubilii Qadri 1969 H. waltairensis Narasimhamurti | |
| 40. | 41. | | *42. | *43. | *44. | *45. | | 46. 47. | 48. | 49. | 50. | 52. | 53. | |

while 2. (Contd.)

| | | 8 | 3 | 4 |
|------------|--|--------------------------------|----------------------------|--|
| | | Genus: Thelohanellus Kudo 1933 | 1933 | |
| 55. | T. rohitae Southwell and | Labeo rohita | Gills | $30-32 \times 7-8$ |
| 56. | Prashad 1918 T. seni Southwell and Prashad 1918 | Catla catla Labeo rohita | Branchiae | $13 \cdot 2 - 13 \cdot 6 \times 10 \cdot 1 - 10 \cdot 3$ |
| 57. | T. catlae Chakravarty and Basu 1948 | Catla catla | Gills | $19.26-21.40 \times 10.7-12.4$ |
| 58. | T. calbasui Tripathi 1952 | Labeo calbasu | Scales | $9-10.8 \times 7.2-5.5$ |
| 59. | T. gangeticus Tripathi 1952 | Labeo rohita, Chela bacaila | Pectoral region | $16.2 - 17.5 \times 5.4 - 3.5$ |
| 90. | T. mrigalae Tripathi 1952 | Cirrhina mrigala | Cysts on head between eyes | $10.8-12 \times 6.3-7.2 \times 4.5-5.4$ |
| 61. | T. andhrae Qadri 1962 | Labeo fimbriatus | Gills | $11.25-14.5 \times 4.5-5.5$ |
| 62. | T. boggoti Qadri 1962 | Labeo boggot | Gills | $11-12 \times 6-7.5$ |
| 63. | T. batae Lalitakumari 1969 | Labeo bata | Gills | $11.4 - 13.6 \times 4.3 - 7.9$ |
| 2 | T. chelae Lalitakumari 1969 | Chela becaila | Bile | $9.4-10.3 \times 4.6-6$ |
| 5 | L. notaili Lalitakumari 1969 | Labeo potail | Gill contents | $12 \cdot 1 - 16 \times 7 \cdot 9 - 9 \cdot 5$ |
| .99 | T. shortii Qadri 1969 | Labeo fibimbriatus | Fins | $11.42-12.85 (12.53) \times 6.42-7.14$ (6.91) |
| | | Genus: Phlogospora Qadri 1962 | 1962 | |
| .19 | P. mysti Qadri 1962 | Mystus bleekeri | Gill contents | 14-18 × 3·5-5 |
| | | Genus: Myxobiatus Davis 1944 | vis 1944 | |
| 8 8 | M. mastacembeli Qadri and Lalitakumari 1965 | Mastacembelus armatus | Gills, intestine | $23.5-30.5 \times 4.6-6.16$ |

| | $79.20-93.6 \times 5.4$ | | 14.1×3.6 | $12-15 \times 8 \cdot 5-10$ $14 \cdot 42 \times 6 \cdot 18$ | $23-27\times 6\cdot 18$ | $11.4-13.6 \times 5.4-7.1$ 15.48×7.7 | | 12.36×6.18 $10.3 \times 4.12 \times 5.18$ $12.9-14.3 \times 5-7.1$ | $9 \cdot 3 - 12 \cdot 9 \times 4 \cdot 3 - 7 \cdot 1$ | | $28.8-30 \times 5-5.5$ 19.8×5.4 |
|-----------------------------------|-------------------------------|--|---|--|---|---|----------------------------------|--|---|-----------------------------------|---|
| thi 1952 | Gills | 1882 | Gall bladder | Gall bladder Gall bladder | Gall bladder | Gall bladder Gut | h 1970 | Gall bladder Gall bladder Gall bladder | Gall bladder | han 1892 | Gall bladder Gall bladder |
| Genus: Neohenneguya Tripathi 1952 | Odontamblyopus rubicundus | Order: Bipolarina Family: Myxidiidae Genus: Myxidiium Butschlii 1882 | Anabas testudineus, Roloonhthalmus hoddaerti | Glossogobius giuris Heteropneustes fossilis | Lates calcarifer | Macrones aor Boleophthalmus boddaerti | Genus: Zschokkella Auerbach 1970 | Hilsa ilisha Heteropneustes fossilis Ophicephalus striatus | gri | Genus: Sphaerospora Thelohan 1892 | Odontamblyopus rubicundus Therapon jarbua |
| | N. tetraradiata Tripathi 1952 | | M. lieberkuhni Buetschlii 1881 | M. glossogobii Chakravarty 1939 M. heteropneustii Chakravarty | M. procerum var. calcarifory Chakravartv 1943 | M. aori Lalitakumari 1969 M. boddaerti Choudhury and Nandi 1973 | | Z. ilishae Chakravarty 1943Z. fossilae Chakravarty 1943Z. ophicephali Lalitakumari | Z. labeonis Lalitakumari 1969 | | S. puttai Tripathi 1952 S. theraponi Tripathi 1952 |
| | .69 | | 70. | 71. | 73. | 74. 75. | | 76. 77. 78. | 13. | | 80. 81. |

Table 3. Myxosporidia reported from Channa (Ophicephalus).

| | Parasite | Host | Site of infection | Description of the (spore Measurements in microns) |
|----|--|--|--|--|
| - | Myxosoma andhrae Lalitakumari 1969 | Ophicephalus punctatus | Outer wall of intestine | Spores pyriform ellipsoidal, in some, the posterior end is drawn into an appendage 1·4-3·6 in length. Spores 12·1-15·7 × 5·7-8·6. Polar capsules unequal. Polar filament length not given. |
| 7 | M. channai n. sp. (present record) | Channa punctata (=Ophicephalus punctatus) | Kidney, Liver, body muscles and fins | Spores pyriform or oval $14.5-18.0 \times 6.0-6.5$. Thick median sutural ridge. Polar capsules dissimilar. Large polar filament $35.0-42.0$, small $20.0-28.0$. No iodinophilous vacuole. |
| œ. | Myxobolus ophicephali Bhatt and Siddiqui 1964 | Ophicephalus punctata | Accessory respiratory organs | Spores pyriform. $11.6-13.3 \times 4.6-6.3$. Polar capsules unequal. Large $6.9-8.5 \times 1.2-2$; Small $6.7-7.3 \times 1.2-2$. |
| 4. | M. aligarhensis Bhatt and Siddiqui 1964 | Channa punctata (= Ophicephalus punctatus) | Accessory respiratory organs | Spores pyriform, shell valves smooth and symmetrical $11\cdot4-15\times6-7\cdot92$. Polar capsule equal $7\cdot6-9\cdot2\times1\cdot2-2\cdot2$ polar filament not mentioned. |
| 5. | M. punctatus Chaudhuri and Chakravarty 1970 | Channa punctata (= Ophicephalus punctatus) | Spleen and pharyngeal epithelium | Spores pyriform $12 \cdot 29 - 15 \times 5 \cdot 72 - 7 \cdot 86$. Pola capsules equal $8 \cdot 57 - 10 \times 2 \cdot 14 - 2 \cdot 86$ Polar firlament $42 \cdot 25 - 32 \cdot 56$. |

| Gills Spores pyriform $19\cdot8-26\cdot4\times5\cdot4-7\cdot2$. Caudal prolongation $20\cdot5-23\cdot8$ polar capsules unequal large: $7\cdot2-8\cdot1\times2-3$ small: $5\cdot5-7\cdot2\times2-3$. Polar filament not recorded. | Mouth cavity Spores oval the posterior end of shell. Pharyngeal valve prolonged into a more or less extended process to form a single caudal appendage. Spore 25·6-39·2 × 3·2-4·4. Caudal appendage 10-20·3. | Branchial epithe- Spore oblongate. Posterior portion gradulum and Gills ally tapering and continued into long caudal prolongation. Characteristically forked at the posterior end. 35·73-45·3 × 2·86-4·29. Caudal prolongation 14·29-27·15. | Roof of buccal Spores oblongate, spore body continued cavity into a long caudal prolongation which is single and undivided. Spore 26.4–29.15 × 2.75–3.85. Caudal prolongation 11.55–13.75. Polar capsules equal, 3.33–3.56 × 1.1–1.38. Polar filament 18–27. | Branchiae and Spore ovoid or oblongate tail bifurcated muscles $41 \cdot 5 - 52 \cdot 5 \times 6 \cdot 18 - 7 \cdot 21$. Caudal prolongation 20–32. Polar capsule $6 \cdot 18 - 9 \cdot 27 \times 2 \cdot 06 - 3$. Polar filament $26 - 32$. | Viscera near Spores oval, caudal appendage bifurcated spleen spore $15 \cdot 4 - 26 \cdot 9 \times 4 \cdot 62 - 5 \cdot 39$. Caudal appendage $6 \cdot 16 - 14 \cdot 63$. Polar capsules unequal. Large: $4 \cdot 62 - 6 \cdot 16 \times 1 \cdot 15 - 1 \cdot 92$ small: $3 \cdot 85 - 4 \cdot 62 \times 1 \cdot 15 - 1 \cdot 92$. Polar filament large $41 \cdot 85 - 53 \cdot 13$, small $28 - 46 \cdot 2$. |
|--|--|---|--|---|--|
| Ophicephalus punctatus and O. gachua | Ophicephalus punctatus | Ophicephalus punctatus | Ophicephalus punctatus | Ophicephalus punctatus | O. gachua |
| Unicauda ophicephali Tripathi 1952 | <i>U. basirii</i> Tripathi 1952 | U. bicornuata Chaudhuri and Chakravarty 1970 | Unicauda bengalensis Chaudhuri and Chakravarty 1970 | Henneguya ophicephali Chakravarty 1939 | H. <i>qadrii</i> Lalitakumari 1965 |
| *6. | * | & * | 6* | 10. | # 7 |

Table 3. (Contd.)

| - | | | | |
|----------|---|------------------------|-----------------------------|---|
| _ | 2 | æ | 4 | 5 |
| 12. | 12. H. zahoori Bhatt and Siddiqui 1964 O. punctatus | O. punctatus | Tissue of gill filaments | Spore biconvex $20-30.6 \times 2.1-3$ caudal appendage bifurcated $12-18.6$. Polar capsules equal $4.9-6.7 \times 0.7-1.1$. |
| 13. | H. jubili Qadri 1969 | Notopterus notopterus | | Only a short description. No measurements given. |
| 4 | H. ganapatiae Qadri 1970 | Notopterus notopterus | Gill contents | Length of the spore 9·28-9·99 (9·72). Breadth of the spore 4·0-4·28 (4·50) Length of the caudal appendage 21-25 (22·25), Length of polar capsule 3·21-3·57 (3·32). Breadth of polar capsule |
| 15. | H. waltairensis Narasimhamurti 1975 | Ophicephalus punctatus | Gill filaments | Spore $14 \cdot 6 - 16 \cdot 5 \times 3 \cdot 2 - 4$ Caudal prolongation $40 - 50$. Polar capsules equal $10 - 12 \times 1 \cdot 6 - 2 \cdot 5$. |
| 16. | Zschokkella ophicephali Lalitakumari 1969 | O. striatus | Gall bladder | Spores gibbous shaped in front view 12.9–14.3 \times 5-7.1. Polar capsule spherical with prominent neck. $3.6-3.9$. Polar filament $61-78$ sporoplasm trapezoid. |

nisms with unbranched tails which are not extensions of the spore valves. In general the anterior end of the tail has a concavity and the spore sits in this cup". Hence he says *U. basirii* Bhatt and Siddiqui is an example of misuse of the definition of Davis—and all the 4 species described above do not fit into the genus. He, however, did not suggest a new place for the above species in any one of the described genera. * Wyatt (1979) while discussing the characters of the genus Unicauda stated that "Unicauda was established by Davis for orga-

thickened at the poles. The 2 polar capsules in M. clarii are pear-shaped measuring $6.18 \times 3.09 \,\mu\mathrm{m}$ with the anterior end drawn out into a narrow tube and open to the outside at the same point (figures 60 and 65 of Chakravarty 1943). The sporoplasm occupies the entire extra capsular region and the 2 nuclei are closely associated (Figs. 63 and 64 of Chakravarty 1943) while in the present form the 2 oval polar capsules are smaller measuring $5.0-6.0 \times 2.0-2.5 \,\mu\mathrm{m}$ and open to the outside independently, one on either side of the median line and the openings are covered by a cushion-like thickening. Further the sporoplasmic nuclei are always present one on either side of the vacuole.

In view of the differences mentioned above, we consider the present form as a new species for which the name Myxobolus tripathii n. sp. is proposed.

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